Pfizer and Moderna COVID jabs found to be contaminated with short DNA fragments, making them potentially TUMORIGENIC and GENOTOXIC

11/15/2023 // Lance D Johnson // Views

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A team of researchers from Ontario, Canada recently discovered the presence of DNA fragments in the monovalent and bivalent Pfizer/BioNTech and Moderna mRNA COVID-19 vaccines. All the vaccines in this study contained billions to hundreds of billions of DNA

fragments, and they each exceeded the DNA adulteration standard of 10 nano-grams per dose set by the *Food and Drug Administration* (FDA) and the *World Health Organization* (WHO). In fact, these batches exceeded FDA standards by 188 to 509 times that limit.

Some of the DNA sequences were highly active promoter/enhancer DNA sequences, and some of the DNA fragments were derived from the SV40 virus. This notorious tumor-causing polyomavirus is derived from monkeys. SV40 was once widely administered to human populations through contaminated polio vaccines, which were produced in naturally-infected SV40 monkey cells.

Moderna once cautioned about insertional mutagenesis from plasmid DNA contamination of mRNA vaccines but couldn't prove theirs weren't contaminated

The mRNA vaccines are highly contaminated with DNA fragments that were never removed during the manufacturing process. These serious bioethics violations were recently discussed by a group of international scientists at the World Council for Health. The expert panel discussed how bacterial DNA from microscopic plasmids remain in the vaccines, long after it multiplies DNA during the mRNA manufacturing process.

Moderna scientists acknowledged the risk of DNA contamination and genotoxicity in their patent for "HPIV3 Vaccines." In the patent application, Moderna wrote:

> Deoxyribonucleic acid (DNA) vaccination is one technique used to stimulate humoral and cellular immune responses to foreign antigens, such as hMPV antigens and/or PIV antigens and/or RSV antigens. The direct injection of genetically engineered DNA (e.g., naked plasmid DNA) into a living host results in a small number of its cells directly producing an antigen, resulting in a protective immunological response. With this technique, however, comes potential problems, **including the possibility of insertional mutagenesis, which could lead to the activation of oncogenes or the inhibition of tumor suppressor genes**.

Here, Moderna admits that "naked" plasmid DNA contamination can put patients at risk of insertional mutagenesis and cancer risk through the inhibition of tumor suppressor genes and the activation of oncogenes. However, Moderna never discloses in their patent application that they were unable to prove that their manufacturing processes for mRNA vaccines are without contamination of plasmid DNA.

FDA, Health Canada and EMA ignore tumorigenic and genotoxicity risk from COVID "vaccines"

This short DNA adulteration is of particular concern due to the potential for insertional mutagenesis and integration of foreign DNA into the human genome. In other words, the COVID vaccines are potentially genotoxic, and the regulatory agencies around the world are ignoring this fact, which has legal and ethical implications.

In fact, the FDA, *Health Canada* and the *European Medicines Agency* (EMA)have all acknowledged that plasmid DNA adulteration does occur, but all these agencies refuse to enforce labeling requirements for DNA fragments, effectively forfeiting informed consent and deceiving public health officials, physicians and patients.

Previous FDA guidance cited the potential for COVID vaccines to contain highly active regulatory sequences of DNA that could lead to insertional mutagenesis (integration). However, this guidance has been ignored and an adulterated product continues to receive FDA approval.

The FDA has even denied that this adulteration exists and refuses to acknowledge the risk that this contamination poses to human health. In response to formal questioning, the FDA issued a statement, declaring that "no safety concerns related to the sequence of, or amount of, residual DNA have been identified." The FDA refuses to take this issue seriously, violating 21 U.S. Code § 351 (Adulterated drugs and devices). The FDA also disregards its own guidance in CPG Sec. 420.100 Adulteration of Drugs Under Section 501(b) and 501(c) of the Act. (Direct Reference Seizure Authority for Adulterated Drugs Under Section 501(b)). Here, the FDA clearly breaches all bio-ethical standards and legal safeguards.

Under Operation Warp Speed, safe and efficacious treatments were suppressed to achieve FDA approval for mRNA experimental gene therapy products. These mRNA experiments were branded as "vaccines" and did not undergo genotoxicity and insertional mutagenesis studies. These studies would have been required prior to use in humans if the product was properly codified and labeled.

The risk for insertional mutagenesis is no laughing matter. If a highly contaminated vaccine initiates this process in the individual, it can lead to cancer of the stem and somatic cells. Combined with lipid nanoparticles, this DNA contamination could cross the placenta and concentrate in ovarian tissue, increasing the likelihood of birth defects. These are just the beginning of the horrors unleashed by this genetic experiment.

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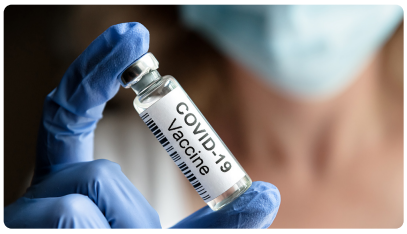

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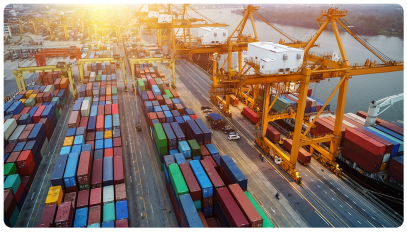

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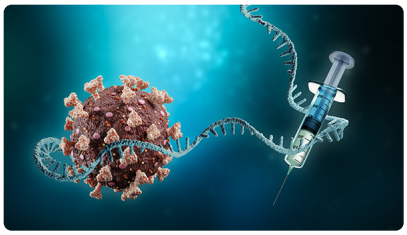

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